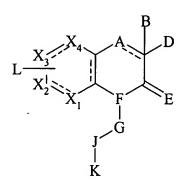
I



or stereoisomers or pharmaceutically acceptable salts, esters, or amides, wherein:

A is selected from NCH<sub>2</sub>, N(alkyl)CH<sub>2</sub>, CH<sub>2</sub>N, CH<sub>2</sub>N(alkyl);

B is selected from H,  $(C_{3-20})$ alkyl, cycloalkyl, heteroalkyl, cycloalkylalkyl, heterocycle, heterocycloalkyl, each optionally substituted with  $R_1$  and  $R_2$ ;

D is selected from H,  $(C_{3-20})$ alkyl, cycloalkyl, heteroalkyl, cycloalkylalkyl, heterocycle, heterocycloalkyl, each optionally substituted with  $R_1$  and  $R_2$ ;

E is absent or selected from O, S, NH;

F is selected from N, NCH<sub>2</sub>, CH<sub>2</sub>N;

G is absent or selected from alkyl, alkyl interrupted by one or more heteroatoms, cycloalkyl, cycloalkyl interrupted by one or more heteroatoms;

J is absent or selected from aryl or heterocycle each optionally substituted with  $R_1$  and  $R_2$ ;

K is absent or selected from an alkyl, alkyl interrupted by one or more heteroatoms, cycloalkyl interrupted by one or more heteroatoms, cycloalkylalkyl interrupted by one or more heteroatoms, each optionally substituted with R<sub>1</sub> and R<sub>2</sub>;

L is selected from H, chlorine, fluorine, bromine, iodine, OH, O(alkyl), amine, alkyl, fluoroalkyl, amide, NO<sub>2</sub>, SH, S(O)<sub>n</sub>(alkyl), SO<sub>3</sub>H, SO<sub>3</sub>alkyl, aldehyde, ketone, acid, ester, urea, Oalkylamide, Oalkylester, Oalkylacid, Nalkylacid, alkylamine, alkylamide, alkylketone, alkylacid, alkylester, alkylurea, Nalkylamide,



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